

(19) World Intellectual Property Organization
International Bureau



(43) International Publication Date
20 September 2001 (20.09.2001)

PCT

(10) International Publication Number
WO 01/68085 A1

(51) International Patent Classification⁷: **A61K 31/195**,
31/185, 31/19, 31/22

(74) Agent: **CARROLL, Lawrence**; Morgan, Lewis & Bock-
ius LLP, 1800 M Street, NW, Washington, DC 20036 (US).

(21) International Application Number: **PCT/US01/08197**

(22) International Filing Date: 15 March 2001 (15.03.2001)

(25) Filing Language: English

(26) Publication Language: English

(30) Priority Data:
60/189,702 15 March 2000 (15.03.2000) US

(71) Applicant (for all designated States except US):
GENAERA CORPORATION [US/US]; 5110 Cam-
pus Drive, Plymouth Meeting, PA 19462 (US).

(72) Inventors; and

(75) Inventors/Applicants (for US only): **FEHLBAUM, Pas-
cale** [FR/US]; 5110 Campus Drive, Plymouth Meeting, PA
19462 (US). **ANDERSON, Mark** [US/US]; 5110 Campus
Drive, Plymouth Meeting, PA 19462 (US). **RAO, Meena**
[IN/US]; 5110 Campus Drive, Plymouth Meeting, PA
19462 (US). **ZASLOFF, Michael** [US/US]; 5110 Campus
Drive, Plymouth Meeting, PA 19462 (US).

(81) Designated States (*national*): AE, AG, AL, AM, AT, AU,
AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU,
CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM,
HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK,
LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX,
MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL,
TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW.

(84) Designated States (*regional*): ARIPO patent (GH, GM,
KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW), Eurasian
patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European
patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE,
IT, LU, MC, NL, PT, SE, TR), OAPI patent (BF, BJ, CF,
CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).

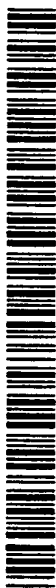
Published:

— with international search report

For two-letter codes and other abbreviations, refer to the "Guid-
ance Notes on Codes and Abbreviations" appearing at the begin-
ning of each regular issue of the PCT Gazette.

(54) Title: A METHOD FOR STIMULATION OF DEFENSIN PRODUCTION

(57) Abstract: The subject invention relates to a method for the stimulation of defensin production in eukaryotic cells such as, for example, mammalian cells. Furthermore, the invention includes said method for the prevention and treatment of infections and other various disease states and in the stimulation of the immune system.



WO 01/68085 A1

TITLE: A METHOD FOR STIMULATION OF DEFENSIN PRODUCTION

Inventors: Pascale Fehlbaum, Mark Anderson, Meena Rao and Michael Zasloff

5

CROSS REFERENCE TO RELATED APPLICATIONS

This application is related to U.S. Patent Application Ser. No.: 09/316,386, filed May 21, 1999, which is herein incorporated by reference in its entirety. This application claims priority to U.S. provisional application 60/189,702, filed March 15, 2000, which is specifically incorporated herein by reference in its entirety.

10

FIELD OF THE INVENTION

This invention relates to stimulating the production of defensins in mammalian cells using the amino acid isoleucine or active isomers or analogs thereof. Furthermore, the present invention includes the use of isoleucine or active isomers or analogs thereof to stimulate defensins for the prevention and treatment of infections and other various disease states.

15

BACKGROUND OF THE INVENTION

Defensins are cationic, cysteine-rich peptides that display broad spectrum antimicrobial activity. Their structure is characterized by a conserved cysteine motif that forms three disulfide linkages, imposing a characteristic β -sheet structure (Hill *et al.*, 1991; White *et al.*, 1995). Associated with this structure is an amphiphilic charge distribution that enables the defensins to interact with and disrupt target cell membranes (Lehrer *et al.*, 1989). This disruption is thought to be accomplished by the formation of channels in the target membrane, leading to cell lysis (Kagan *et al.*, 1990). Defensins have been shown to inhibit proliferation of both gram-positive and gram-negative bacteria, yeast and numerous viruses. In particular, defensins inhibit the proliferation of the yeast strain *Candida albicans* and the gram-negative bacteria *Escherichia coli* (Porter *et al.*, 1997; Harder *et al.*, 1997; Schonwetter *et al.*, 1995; Daher *et al.*, 1986).

20

25

30

Defensins have recently been identified as an integral component of the antimicrobial barrier of mucosal surfaces. In both the human and murine small intestine,

defensin RNA has been localized to the Paneth cell, a specialized epithelial cell located at the crypt base (Ouellette *et al.*, 1989; Jones *et al.*, 1992). The associated peptide has been localized within secretory granules of the Paneth cell and in the lumen of the small intestine, suggesting a role for defensins in host defense in the gut (Selsted *et al.*, 1992).

5 Defensins have also been found in bovine and human respiratory epithelium. Tracheal antimicrobial peptide, a β -defensin isolated from bovine tracheal mucosa, was localized to the ciliated columnar epithelial cells of the trachea and bronchi (Diamond *et al.*, 1991; Diamond *et al.*, 1993). Lingual antimicrobial peptide, another α -defensin, was found in bovine lingual mucosa and stratified squamous epithelium of the tongue (Schonwetter *et al.*, 1995). Most recently, human β -defensin-1 was demonstrated to be present in the epithelium of the trachea and bronchi, as well as the submucosal gland and alveolar epithelium (Goldman *et al.*, 1997; Zhao *et al.*, 1996).

Considerable data exists indicating that epithelial defensins are up-regulated in response to infection. In cultured tracheal epithelial cells, tracheal antimicrobial peptide message is induced following exposure to bacterial lipopolysaccharide (Diamond *et al.*, 15 1996). This induction was blocked by antibody to CD14, suggesting that epithelial cells provide an active, inducible antimicrobial defense. Following injury to bovine tongue, lingual antimicrobial peptide RNA message increased at the site of injury (Schonwetter *et al.*, 1995). Induction of lingual antimicrobial peptide was also observed following acute infection in bronchial epithelium and chronic infection in ileal mucosa (Stolzenberg *et al.*, 20 1997). Together these data support a role for α -defensins as important host defense effector molecules that are rapidly mobilized by epithelium upon injury or infection.

Due to the significant host defense properties of defensins, any means which stimulates or induces the production of these peptides is desired in the art. The present invention provides such means as to stimulate the production of defensins. 25

Some bacteria present in the GI tract of mammals such as Lactobacilli, Bifidobacteria, and Streptococci are considered to be beneficial to the host and are termed probiotic bacteria. Probiotic bacteria are symbiotic organisms in vertebrate hosts and their direct administration or the promotion of their growth is known to prevent and/or treat infection, promote weight gain in animals, and decrease serum lipid levels (Taylor, 30 G.R.J and Williams, 1998; Mohan, B. *et al.*, 1996; Pascual, M. *et al.*, 1999; Hofacre, C.L.

et al., 1998; Vanderhóff, J.A. et al., 1999). Materials with the ability to stimulate the growth of probiotic bacteria are known as prebiotic materials (Collins and Gibson, 1999). Through their effects in promoting probiotic bacterial numbers prebiotic materials are also thought to provide their benefits (Gibson, G.R., 1999; Bovee-Oudenhoven, I.M et al.).

SUMMARY OF THE INVENTION

The present invention comprises a method of increasing the production of defensins in eukaryotic cells. This method comprises exposing the eukaryotic cells to a composition comprising isoleucine or active isomers or analogs thereof in an amount sufficient to effect said increase. The method also comprises increasing defensin production in eukaryotic cells using isomers of isoleucine including stereoisomers, diastereomers in particular or a combination thereof. The stereoisomers include L-isoleucine, D-isoleucine and D-allo-isoleucine. The method further comprises increasing defensin production in eukaryotic cells using active analogs of isoleucine including alpha-keto-methyl-valerate, isoleucine hydroxamate, butyrate, and valine.

The eukaryotic cells where defensin production may be stimulated may be mammalian cells, and more particularly, epithelial cells. These epithelial cells may be from a tissue or source selected from, for example, the group comprising brain, kidney, heart, spleen, buccal mucosa, nasal mucosa, conjunctiva, tongue, choroid plexus, trachea, bronchi, bronchioles, fallopian tubes, uterus, cervix, vagina, testes, bladder, urethra, esophagus, duodenum, jejunum, ileum, caecum, ascending colon, sigmoid colon, descending colon and rectum.

Furthermore, the invention includes a method of treating or preventing an infection or other disease state in a patient. This method comprises administering a composition comprising isoleucine or active isomers or analogs thereof in an amount sufficient to effect treatment or prevention. The infection may be caused by any viral, bacterial or fungal pathogen including, for example, *Candida albicans*, *Escherichia coli*, Rotavirus or Respiratory Syncytial Virus. The invention also encompasses a method of stimulating the immune system of a mammal comprising administering to the mammal,

isoleucine or active isomers or analogs thereof in an amount sufficient to effect the stimulation.

The present invention also includes a method of stimulating the growth of beneficial probiotic bacteria such as Lactobacilli, Bifidobacteria, and Streptococci. The defensin inducing compounds described in the invention are expected to act as prebiotic materials and therefore stimulate the growth of probiotic bacteria.

The present invention also includes a method for improving the general health, total weight gain, rate of weight gain, efficiency of feed conversion, and/or reduction or elimination of carriage of pathogenic organisms in livestock. These benefits are known or are expected benefits of probiotic bacteria and therefore are also expected due to the prebiotic effects described above.

The accompanying figures, which are incorporated in and constitute a part of this specification, illustrate several embodiments of the invention and together with the description, serve to explain the principle of the invention.

BRIEF DESCRIPTION OF THE FIGURES

Figure 1: Effect of various amino acids on defensin production in MDBK cells.

Figure 2: Dose-response effect of L-isoleucine on defensin production in MDBK cells.

Figure 3: Dose-response effect of D-isoleucine on defensin production in MDBK cells.

Figure 4: Comparison of effects of isoleucine and alloisoleucine on defensin production in MDBK cells.

Figure 5: Effect of L-isoleucine on defensin production in the human colon epithelial cell line HT-29.

Figure 6: Defensin inducing effect of alpha-keto-methylvalerate.

Figure 7: Defensin inducing effect of butyrate.

Figure 8: Defensin inducing effect of isoleucine hydroxamate.

Figure 9: Defensin inducing effect of valine.

Figure 10: Illustration of generic structure of defensin inducing isoleucine analogs.

DETAILED DESCRIPTION OF THE INVENTION

Definitions

“Consisting essentially of” herein refers to compositions wherein the named chemical constituent, active isomer or analog thereof is the principal ingredient of said composition.

“Active isomer” herein refer to molecules having the same molecular formula of a named chemical constituent but differing in the nature or sequence of binding of their atoms or in the spatial arrangement of their atoms, wherein said molecules elicit defensin production. In a related aspect, “analogs” herein refer to molecules having the generic or similar structure of a named chemical constituent (e.g., corresponding -keto form of a named amino acid), wherein said “analogs” elicit defensin production.

The subject invention relates to a method of stimulating the production of defensins in eukaryotic cells using a composition comprising isoleucine or active isomers or analogs thereof. The composition used in the method may be used to prevent or treat various disease states or conditions. Consequently, a method of treating or preventing an infection or other disease state in a patient may comprise administering a composition comprising isoleucine or active isomers or analogs thereof, in an amount sufficient to effect the treatment or prevention of said infection or disease state.

As exemplified in Figure 1, L-isoleucine has the ability to stimulate the production of defensins in eukaryotic cells. L-isoleucine was capable of stimulating defensin production by MDBK cells at concentrations as low as three micrograms per milliliter. None of the other similar amino acids tested at this concentration had any effect on defensin production. L-valine and L-tyrosine methyl ester had no effect on defensin production at concentrations as high as fifty micrograms per milliliter. Furthermore, L-isoleucine increased defensin production at concentrations fifty to one-hundred-fold less than those of D-isoleucine, indicating that the desired effect of the method is dependent on the stereochemical configuration of isoleucine. Figure 2 demonstrates the dose-dependent effect of L-isoleucine on defensin production, indicating the specific effect of L-isoleucine. Thus, the present invention encompasses a method of eliciting the production of defensins by eukaryotic cells. This method comprises exposing the cells to

the composition containing isoleucine or active isomers or analogs thereof in an amount sufficient to elicit the production of defensins by the cells.

In one embodiment, isoleucine or active isomers or analogs are exposed to cells *ex vivo* at concentrations which elicit the highest production of defensin. Preferably, cells
5 are exposed to concentrations of L-isoleucine from about 3.12 $\mu\text{g/ml}$ to about 100 $\mu\text{g/ml}$, more preferably from about 6.25 $\mu\text{g/ml}$ to about 50 $\mu\text{g/ml}$, even more preferably from about 6.25 $\mu\text{g/ml}$ to about 25 $\mu\text{g/ml}$. Further, cells can be exposed to D-isoleucine at concentrations which elicit the production of defensin. Preferably, cells are exposed to concentrations of D-isoleucine from about 100 $\mu\text{g/ml}$ to about 400 $\mu\text{g/ml}$, more
10 preferably from about 200 $\mu\text{g/ml}$ to about 400 $\mu\text{g/ml}$.

The method of the invention encompasses a composition comprising isoleucine or active isomers thereof. Isomers of isoleucine comprise both stereoisomers and diastereomers as isoleucine has two chiral centers allowing for four separate stereoisomers. The importance of stereochemistry to the present invention will be
15 apparent to one skilled in the art because L-isoleucine stimulated defensin production at concentrations approximately fifty to one-hundred-fold less than D-isoleucine as exemplified in Figures 2 and 3. Similarly, changing the configuration of isoleucine at its second chiral center yields the compound alloisoleucine. Figure 4 shows that alloisoleucine is not as effective an inducer of defensin production in MDBK cells as is
20 isoleucine. The strong dependence of defensin inducing activity on the chiral configuration of isoleucine supports the specificity of isoleucine as a defensin inducer. Thus, the present method of the invention comprises a method whereby the composition for stimulating defensin production contains L-isoleucine, D-isoleucine, D-alloisoleucine, or a mixture thereof.

25 Figure 5 illustrates the defensin inducing property of L-isoleucine in the human colon epithelial cell line HT-29. This result, taken together with similar data from MDBK cells shows that isoleucine has utility as a defensin inducer in a variety of species and at a variety of epithelial surfaces that may be of therapeutic importance.

In addition to L-isoleucine and D-isoleucine other similar molecules also act as
30 defensin inducers. Figures 6, 7, and 8 demonstrate that the compounds alpha-keto-methylvalerate, butyrate, and isoleucine hydroxamate are inducers of defensin production

in epithelial cells. Figure 9 shows that valine stimulates defensin expression at concentrations that are substantially higher than those needed for isoleucine but which may have utility in treating or preventing infection. In a preferred embodiment, Figure 10 illustrates a generic structure for defensin inducing isoleucine analogs.

5 The method of the present invention comprises stimulation of defensin production by epithelial cells derived from, for example, the following mammalian tissues: brain, skin, kidney, heart, spleen, buccal mucosa, nasal mucosa, conjunctiva, tongue, choroid plexus, trachea, bronchi, bronchioles, fallopian tubes, uterus, cervix, vagina, testes, bladder, urethra, esophagus, duodenum, jejunum, ileum, caecum, ascending colon,
10 sigmoid colon, descending colon and rectum. The method may also be used to stimulate defensin production in epithelial cells found in other tissues, for example, the ear, liver, pancreas or ovary. For example, the method may be utilized for the treatment or prevention of dermal, oral, ocular, respiratory, gastrointestinal, colorectal and urogenitary diseases or other epithelial cell-related diseases in mammals, including humans and
15 animals.

 The method of the invention is useful in treating or preventing infections resulting from a broad range of pathogens as defensins have been shown to inhibit proliferation of both gram-positive and gram-negative bacteria, yeast and numerous viruses. For example, the method is effective in the treatment of candidiasis because defensins inhibit
20 the proliferation of the underlying pathogen *Candida albicans*. The method is also be useful in treating diarrhea, dysentery, septicemia and acute infantile gastroenteritis as defensins are known to inhibit proliferation of the underlying pathogens of these disease states, particularly *Escherichia coli*. Treatment of acute respiratory disease resulting from Respiratory Syncytial Virus will also be possible as defensins inhibit the proliferation of
25 such viruses.

 The present invention also includes a method of stimulating the immune system of a mammal after, for example, surgery, immune ablation by chemotherapy or other treatments, or bacterial or viral infections. Such a method comprises administering a composition comprising isoleucine or active isomers or analogs thereof to the patient,
30 human or animal, requiring immune system stimulation in an amount sufficient to effect such stimulation. Thus, stimulation of defensin production in the epithelial cells of the

patient is the mechanism whereby stimulation of the immune system occurs. Further, methods are envisaged which comprise immunostimulation by administering compositions comprising isoleucine or active isomers or analogs thereof and cytokines or other immune stimulants. Administration of such compositions are preferred for diseases involving gram-negative infections and septic conditions wherein LPS (lipopolysaccharide) is the major immunogenic agent. In a preferred embodiment, administration can be sequential or more preferably simultaneous (i.e., co-administered).

The present invention also includes a method of stimulating the growth of beneficial probiotic bacteria such as Lactobacilli, Bifidobacteria, and Streptococci. Materials with this property are known as prebiotic materials (Collins and Gibson, 1999). The defensin inducing compounds described in the invention are expected to act as prebiotic materials. Probiotic bacteria are relatively much less sensitive to the killing effects of antimicrobial peptides. Induction of defensin or other antimicrobial peptide production at an epithelial surface is expected to lead to direct killing of sensitive, non-probiotic bacteria including pathogens while having little direct effect on the relatively insensitive probiotic bacteria. This is expected to provide a competitive advantage to the probiotic bacteria in the internal microbial ecology of the treated host as they gain nutrients and space at the expense of the sensitive microbes. It is known that heat killed probiotic bacteria are themselves inducers of epithelial defensins and that probiotic bacteria provide benefit to the host; this relationship is then truly symbiotic in that it provides benefits to both host and probiotic bacteria. This effect can be utilized to promote probiotic bacterial numbers and thereby provide the preventive and therapeutic benefits described above. The prebiotic effect of defensin inducers may be useful in at any epithelial surface that is able to support probiotic or commensal bacteria, especially the entire digestive tract, upper airway, and vagina.

The present invention also includes a method for improving the general health, total weight gain, rate of weight gain, efficiency of feed conversion, and/or reduction or elimination of carriage of pathogenic organisms in livestock. These benefits are known or are expected benefits of probiotic bacteria and therefore are also expected due to the prebiotic effects described above. The direct microbicidal effect of defensins induced in treated animals is also expected to mediate these effects by killing pathogens and

improving the general health of the animals, allowing for improved average weight gain as less energy is expended by the animals in fighting pathogens.

The present invention also includes a method for increasing numbers of probiotic or commensal bacteria comprising the administration of isoleucine or active isomers thereof combined with live probiotic or commensal bacteria and/or other prebiotic materials including non-digestible oligosaccharides such as fructo-oligosaccharides, inulin, or chicory.

The invention includes pharmaceutical compositions comprising isoleucine or active isomers or analogs thereof or a combination of isoleucine or active isomers or analogs thereof, together with a pharmaceutically acceptable carrier. Preferred embodiments may include such compositions comprising purified isoleucine, active isomers or analogs or combinations of any of the mentioned compounds. Acceptable carriers can be sterile liquids, such as water and oils, including those of petroleum, animal, vegetable or synthetic origin, such as peanut oil, soybean oil, mineral oil, sesame oil and the like. Water is a preferred carrier when the pharmaceutical composition is administered intravenously. Saline solutions and aqueous dextrose and glycerol solutions can also be employed as liquid carriers, particularly for injectable solutions. Suitable pharmaceutical carriers are described in Remington's Pharmaceutical Sciences, 19th edition, Mack Publishing Company, 1995. The pharmaceutical compositions used in the method of treatment of this invention may be administered systemically or topically, depending on such considerations as the condition to be treated, need for site-specific treatment, quantity to be administered and similar considerations. This administration may be by the oral, intravenous, or inhaled route or by suppository, enema, mouth wash or the like.

Topical administration may be used. Any common topical formation such as a solution, suspension, gel, ointment or salve and the like may be employed. Preparation of such topical formulations are well described in the art of pharmaceutical formulations as exemplified, for example, by Remington's Pharmaceutical Sciences, 19th edition, Mack Publishing Company, 1995. For topical application, these compounds could also be administered as a powder or spray, particularly in aerosol form or as a lozenge for local oral delivery. The active ingredient may be administered in pharmaceutical compositions

adapted for systemic administration. As is known, if a drug is to be administered systemically, it may be confectioned as a powder, pill, tablet or the like or as a syrup or elixir for oral administration. For intravenous, intraperitoneal or intralesional administration, the compound will be prepared as a solution or suspension capable of being administered by injection. In certain cases, it may be useful to formulate these compounds in suppository form or as an extended release formulation for deposit under the skin or intramuscular injection.

The compounds can also be administered in animal feed for the stimulation of weight gain and to enhance animal health.

An effective amount is that amount which will increase the expression of defensins. A given effective amount will vary from condition to condition and in certain instances may vary with the severity of the condition being treated and the patient's susceptibility to treatment. Accordingly, a given effective amount will be best determined at the time and place through routine experimentation as shown in Figure 2. However, it is anticipated that in the treatment and prevention of infections and other disease states is in accordance with the present invention, a formulation containing between 0.001 and 5.0 % by weight, preferably about 0.01 to 1.0 %, will usually constitute an effective therapeutic amount. When administered systemically, an amount between 0.01 and 100 milligrams per kilogram body weight per day, but preferably about 0.1 to 10 milligrams per kilogram, will effect a therapeutic result in most instances.

The present compositions are preferably for treatment of human subjects, however, the mentioned compositions are also contemplated for use in animal subjects, including farm animals as well as domestic species.

The practice of the present invention will employ the conventional terms and techniques of molecular biology, pharmacology, immunology and biochemistry that are within the ordinary skill of those in the art. For example, see Sambrook *et al.*, Molecular Cloning: A Laboratory Manual, 3rd edition, Cold Spring Harbor Laboratory Press, 1989.

Other embodiments of the invention will be apparent to those skilled in the art from consideration of the specification and practice of the invention disclosed. It is intended that the specifications and examples be considered exemplary only with the true

scope of the invention being indicated by the claims. Having provided this detailed information, applicants now describe preferred aspects of the invention.

EXAMPLE 1

Defensin induction in MDBK cells.

Cell Culture: MDBK (Madin-Darby Bovine Kidney) cells were obtained from the ATCC (Rockville, MD) and were maintained in growth medium consisting of Eagle's modified essential media with Earle's balanced salt solution, 10% horse serum, 0.10 mM non-essential amino acids and no antibiotics. For stimulation experiments, cells were
10 plated into six well tissue culture plates and maintained for three days in growth medium until cells were almost confluent. The medium was then changed to serum-free epithelial cell growth medium (Clonetics, San Diego, CA) and the test material was added to the dish. Twenty-four hours later, the medium was withdrawn and cells were rinsed with phosphate-buffered saline. Total RNA was then isolated using Trizol reagent
15 (Gibco/BRL, Grand Island, NY) according to protocols supplied by the manufacturer. RNA was quantified by measuring the OD₂₆₀ of each sample.

RNA-Polymerase Chain Reaction: Total RNA was treated with DNase prior to reverse transcription and PCR. The DNase was heat inactivated at 65 C in the presence of EDTA for ten minutes. Reverse transcription and PCR were performed essentially as
20 described for the GeneAmp RNA PCR kit (Perkin Elmer, Foster City, CA). Briefly, approximately 250 nanograms of total RNA was primed with polydT and reverse transcribed with Murine Leukemia Virus reverse transcriptase in a total volume of 16 µL at room temperature for ten minutes and then at 42 C for an additional fifteen minutes. The reverse transcriptase was heat inactivated at 99 C for five minutes and the reaction
25 was chilled to 4 degrees C. This reverse transcription reaction was then split in half: one portion was used for amplification of the target defensin RNA, and the other was treated in parallel to determine the β-tubulin RNA level as a control. Additional reagents necessary for the PCR reaction, including appropriate synthetic DNA primers;

30 β-defensin 5' CTC TTC CTG GTC CTG TCT 3' (SEQ ID NO: 1)
 5' CTT CTT TTA CTT CCT CCT GCA GCA 3' (SEQ ID NO: 2)

β -tubulin 5' GTT CCC AAA GAT GTC AAT GCT GCC 3' (SEQ ID NO: 3)

5' ATG CTG CAA GGC TGA AAG GAA TGG 3' (SEQ ID NO: 4)

5 were added after splitting the reverse transcription reactions to bring the reaction volumes to 40 μ L. The reactions were then subjected to thermal cycling as follows: 95 C for one minute, 52 C for one minute, 72 C for one minute for 30 cycles followed by a single 72 C incubation for fifteen minutes to allow for extension. The expected 200 base pair β -defensin PCR product was measured by gel electrophoresis or QPCR (Quantitative
10 PCR).

Product Capture for QPCR System: 10 μ L of the final PCR mixture was combined with 15 μ L of strepavidin bead slurry (Perkin-Elmer, Foster City, CA), 21 μ L H₂O and 4 μ L of 10x PCR buffer to yield a 50 μ L binding reaction. The binding reactions were incubated at room temperature for fifteen minutes with occasional agitation. One ml of
15 QPCR assay buffer was then added to the reaction. The quantity of PCR product was subsequently measured with the Perkin-Elmer QPCR instrument (Perkin-Elmer, Foster City, CA). The amount of defensin product was normalized to endogenous expression in MDBK cells in the absence of any experimental agent.

EXAMPLE 2

20 Defensin induction in MDBK cells.

In order to easily identify compounds that have defensin inducing activity a stable cell line containing an integrated plasmid in which expression of the easily assayed gene product luciferase is controlled by a bovine beta-defensin promoter was constructed.
Cloning of bovine defensin promoter: A DNA fragment containing the bovine enteric
25 beta-defensin (EBD) promoter was generated via PCR. The fragment contained 812 base pairs of 5'-flanking sequence and the first 43 base pairs of the 5' untranslated portion of the EBD cDNA. This DNA fragment was engineered to contain an Mlu I restriction site at the 5' end of the fragment and a Bgl II restriction site at the 3' end to facilitate subsequent cloning into the pGL2-basic luciferase expression plasmid (Promega). The PCR product
30 was cloned into the TA cloning vector (Invitrogen) by standard techniques and sequenced to confirm its identity.

Construction of EBD promoter-luciferase reporter plasmid: The defensin promoter containing TA-vector plasmid was digested with Mlu I and Bgl II and the appropriate digestion product was isolated following separation on a 1.2% agarose gel. The luciferase expression vector pGL2-basic was similarly digested with Mlu I and Bgl II and isolated following gel electrophoresis. The vector and defensin promoter fragment were ligated together and transformed into *E. coli* via standard procedures.

Generation of Stable MDBK cell lines containing a functional defensin promoter-luciferase plasmid: The defensin promoter-luciferase plasmid was mixed with the G418 resistance plasmid LNCZ in a 1 to 5 ratio, combined with Fugene™ transfection reagent and placed on MDBK cells.

Cells were then exposed to medium containing 0.4 milligrams/ml G418 until resistant colonies were visible (4-5 weeks). The resistant clones were then expanded and screened for the expression of luciferase.

Screening of compounds for defensin inducing activity: Cells of a clonal MDBK cell line expressing the defensin promoter-luciferase plasmid were placed in the wells of a 96-well tissue culture plate. Test substances were then placed in tissue culture medium and added to the wells. After the test substances had been in contact with the cells for 12-24 hours luciferase expression levels were measured using standard procedures.

20

EXAMPLE 3

Stimulation of probacterial growth and weight gain in domestic animals

Isoleucine is added to the feed of livestock in amounts varying from .0025 to .05 Wt% and these animals are compared with animals fed unsupplemented feed or feed with added antibiotics. Animals are weighed and blood samples are drawn weekly. Weight values are used to determine total weight gain, rate of weight gain and feed conversion efficiency. Blood samples are tested for levels of circulating IgA, IgG, IgM and isoleucine. Fecal samples are also obtained weekly and tested for numbers of cultured micro-organisms/gram of shed feces (Identification and quantification of species of interest, including Lactobacilli, Streptococci, Bifidobacteria, *E. coli*, Bacteriodes, Salmonella, Clostridia, total aerobes, and total anaerobes). Levels of fecal immunoglobulins are also determined.

While the invention has been described and illustrated herein by references to various specific materials, procedures and examples, it is understood that the invention is not restricted to the particular combinations of material and procedures selected for that purpose. Numerous variations of such details can be implied as will be appreciated by those skilled in the art. All references, patents or other publications cited in this application or in the following reference section are herein incorporated by reference in their entirety.

REFERENCES

- Bovee-Oudenhoven, IM, Wissink, ML, Wouters, JT, and Van der Meer, R. *Dietary calcium phosphate stimulates intestinal lactobacilli and decreases the severity of a salmonella infection in rats.* J Nutr. 129, 607-12, 1999.
- 5 Collins MD and Gibson GR. *Probiotics, prebiotics, and synbiotics: approaches for modulating the microbial ecology of the gut.* Am. J. Clin. Nutr. 69 (suppl), 1052S-1057S, 1999.
- Daher KA, Selsted ME and Lehrer RI. *Direct inactivation of viruses by human granulocyte defensins.* J. Virol. 60, 1068-1074, 1986.
- 10 Diamond G, Zasloff M, Eck H, Brasseur M, Maloy WL and Bevins CL. *Tracheal antimicrobial peptide, a cysteine-rich peptide from mammalian tracheal mucosa: peptide isolation and cloning of a cDNA.* Proc. Natl. Acad. Sci. USA 88, 3952-3956, 1991.
- Diamond G, Jones DE and Bevins CL. *Airway epithelial cells are the site of expression of a mammalian antimicrobial peptide gene.* Proc. Natl. Acad. Sci. USA 90, 4596-4600.
- 15 1993.
- Diamond G, Russell JP and Bevins CL. *Inducible expression of an antibiotic peptide gene in lipopolysaccharide-challenged tracheal epithelial cells.* Proc. Natl. Acad. Sci. USA 93, 5156-5160, 1996.
- Gibson, GR. *Dietary modulation of the human gut microflora using the prebiotics oligofructose and inulin.* J Nutr. 129(7 Suppl), 1438S-1441S, 1999.
- 20 Goldman MJ, Anderson GM, Stolzenberg ED, Kari UP, Zasloff M and Wilson JM. *Human beta-defensin-1 is a salt-sensitive antibiotic in lung that is inactivated in cystic fibrosis.* Cell 88, 553-560, 1997.
- Harder J, Bartels J, Christophers E and Schröder JM. *A peptide antibiotic from human skin.* Nature 387, 861, 1997.
- 25 Hill CP, Yee J, Selsted ME and Eisenberg D. *Crystal structure of defensin HNP-3, an amphiphilic dimer: mechanisms of membrane permeabilization.* Science 251, 1481-1485, 1991.
- Hofacre, CL, Froyman, R, Gautrias, B, George, B, Goodwin, MA, and Brown, J. *Use of Aviguard and other intestinal bioproducts in experimental Clostridium perfringens-associated necrotizing enteritis in broiler chickens.* Avian Diseases 42, 579-584, 1998.
- 30

- Jones DE and Bevins CL. *Paneth cells of the human small intestine express an antimicrobial peptide gene*. J. Biol. Chem. 267, 23216-23225, 1992.
- Lehrer RI, Barton A, Daher KA, Harwig SS, Ganz T and Selsted ME. *Interaction of human defensins with Escherichia coli. Mechanism of bactericidal activity*. J. Clin. Invest. 84, 553-561, 1989.
- Mohan, B, Kadirvel, R., Natarajan, A., and Bhakaran, M. *Effect of probiotic supplementation on growth, nitrogen utilization and serum cholesterol in broilers*. British Poultry Science 37, 395-401, 1996.
- Ouellette AJ, Greco RM, James M, Frederick D, Naftilan J and Fallon JT. *Developmental regulation of cryptdin, a corticostatin/defensin precursor mRNA in mouse small intestinal crypt epithelium*. J. Cell Biol. 108, 1687-1695, 1989.
- Pascual, M, Hugas, M, Badiola, JJ, Monfort, JM, and Garriga, M. *Lactobacillus salivarius CTC2197 prevents Salmonella enteritidis colonization in chickens*. Applied and Environmental Microbiology, 65, 4981-4986, 1999.
- Porter EM, Van Dam E, Valore EV and Ganz T. *Broad-spectrum antimicrobial activity of human intestinal defensin five*. Infect. Immun. 65, 2396-2401, 1997.
- Kagan BL, Selsted ME, Ganz T and Lehrer RI. *Antimicrobial defensin peptides form voltage-dependent ion-permeable channels in planar lipid bilayer membranes*. Proc. Natl. Acad. Sci. USA 87, 210-214, 1990.
- Schonwetter BS, Stolzenberg ED and Zasloff M. *Epithelial antibiotics induced at sites of inflammation*. Science 267, 1645-1648, 1995.
- Selsted ME, Miller SI, Henschen AH and Ouellette AJ. *Enteric defensins: antibiotic peptide components of intestinal host defense*. J. Cell Biol. 118, 929-936, 1992.
- Stolzenberg ED, Anderson GM, Ackerman MR, Whitlock RH and Zasloff M. *Epithelial antibiotic induced states of disease*. Proc. Natl. Acad. Sci. USA 94, 8686-8690, 1997.
- Taylor, GRJ and Williams, CM. *Effects of probiotics and prebiotics on blood lipids*. British Journal of Nutrition, 80, Suppl.2, S225-S230, 1998.
- Vanderhoof, JA, Whitney, DB, Antonson, DL, Hanner, TL, Lupo, JV, Young, RJ. *Lactobacillus GG in the prevention of antibiotic-associated diarrhea in children*. Journal of Pediatrics, 135, 564-568, 1999.

White SH, Wimley WC, and Selsted ME. *Structure, function, and membrane integration of defensins*. Curr. Opin Struct. Biol. 5, 521-527, 1995.

Zhao C, Wang I and Lehrer RI. *Widespread expression of beta-defensin hBD-1 in human secretory glands and epithelial cells*. FEBS Lett. 396, 319-322, 1996.

What is claimed:

1. A method of eliciting the production of defensins in eukaryotic cells comprising exposing said cells to a composition comprising isoleucine or
5 active isomers or analogs thereof in an amount sufficient to elicit said production.
2. The method of claim 1, wherein said composition consists essentially of isoleucine or active isomers or analogs thereof in an amount sufficient to elicit said production.
- 10 3. The method of claim 1, wherein said composition further comprises a cytokine.
4. The method of any one of claims 1-3, wherein said eukaryotic cells are
15 mammalian cells.
5. The method of claim 4, wherein said mammalian cells are epithelial cells.
6. The method of claim 5, wherein said epithelial cells are from a tissue or source
20 selected from the group consisting of brain, kidney, heart, spleen, buccal mucosa, nasal mucosa, conjunctiva, tongue, choroid plexus, trachea, bronchi, bronchioles, fallopian tubes, uterus, cervix, vagina, testes, bladder, urethra, esophagus, duodenum, jejunum, ileum, caecum, ascending colon, sigmoid colon, descending colon and rectum.
- 25 7. A method according to any one of claims 1-3, wherein the isomers are stereoisomers.
8. A method according to claim 7, wherein the stereoisomer is L-isoleucine.
- 30 9. A method according to claim 7, wherein the stereoisomer is D-isoleucine.

10. A method according to claim 7, wherein the stereoisomer is D-allo-isoleucine.

5 11. A method according to any one of claims 1-3, wherein the analog is alpha-keto-methylvalerate.

12. A method according to any one of claims 1-3, wherein the analog is isoleucine hydroxamate.

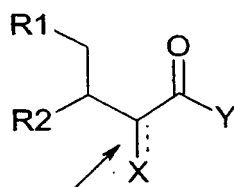
10

13. A method according to any one of claims 1-3, wherein the analog is valine.

14. A method according to any one of claims 1-3, wherein the analog is butyrate or an active derivative thereof.

15

15. A method according to any one of claims 1-4, wherein the analog is one of a member of a class of compounds defined by the chemical structure drawing below:



single or double bond

X = -NH₂, O, H.

Y = -OH, -NHOH, -NH₂, -OCH₃, -O-R₃.

R₁ = H, -CH₃, alkyl of 2-5 carbons.

R₂ = H, -CH₃, alkyl of 2-5 carbons.

R₃ = alkyl of 2-5 carbons.

20

16. A method of treating or preventing an infection or other disease state in a patient in need of said treatment or prevention by administering to said patient a composition consisting essentially of isoleucine or active isomers or analogs thereof in an amount sufficient to effect the treatment or prevention.

25

17. The method of claim 16, wherein said infection is caused by any viral, bacterial or fungal pathogen.

18. The method of claim 13, wherein the pathogen is selected from the group consisting of *Candida albicans*, *Escherichia coli*, Rotavirus or Respiratory Syncytial Virus.

19. A method of stimulating the immune system of a mammal or other animal comprising administering to said mammal or other animal a composition comprising isoleucine or active isomers or analogs thereof in an amount sufficient to effect said stimulation.

20. The method of claim 16, wherein the isoleucine or active isomer or analog thereof is administered after surgery, immune ablation or bacterial or viral infection.

21. A method of increasing numbers of probiotic or commensal bacteria at an epithelial surface or in an epithelially lined compartment comprising administration of a composition comprising isoleucine or active isomers or analogs thereof in an amount sufficient to elicit said increase in bacteria.

22. A method of increasing numbers of probiotic or commensal bacteria at an epithelial surface or in an epithelially lined compartment comprising administration of a composition containing an antimicrobial peptide inducing compound or material.

23. A composition comprising isoleucine or active isomers or analogs thereof in an amount sufficient to elicit increases in probiotic or commensal bacteria.

24. A composition comprising isoleucine or active isomers or analogs thereof and other prebiotic substances.

25. A composition comprising isoleucine or active isomers or analogs thereof and non-digestible carbohydrates including fructo-oligosaccharides (FOS), inulin, and/or chicory singly or in combination.

5

26. A composition comprising isoleucine or active isomers or analogs thereof and live probiotic or commensal bacteria.

10

27. A composition comprising isoleucine or active isomers or analogs thereof and non-digestible carbohydrates including fructo-oligosaccharides (FOS), inulin, and/or chicory singly or in combination and live probiotic bacteria.

15

28. A composition comprising an antimicrobial peptide inducing compound or material and non-digestible carbohydrates including fructo-oligosaccharides (FOS), inulin, and/or chicory singly or in combination.

20

29. A composition comprising an antimicrobial peptide inducing compound or material and live probiotic or commensal bacteria.

30. A composition comprising an antimicrobial peptide inducing compound or material and non-digestible carbohydrates including fructo-oligosaccharides (FOS), inulin, and/or chicory singly or in combination and live probiotic bacteria.

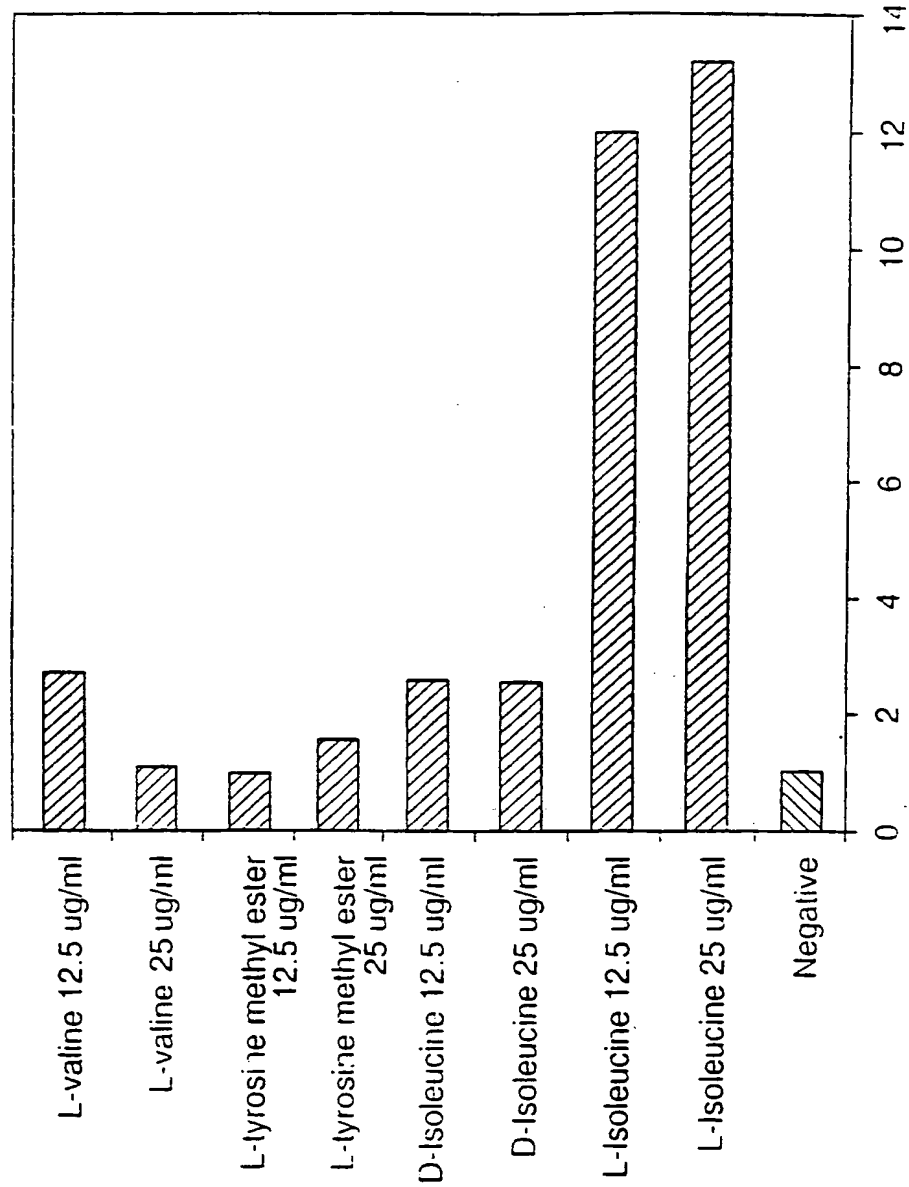
25

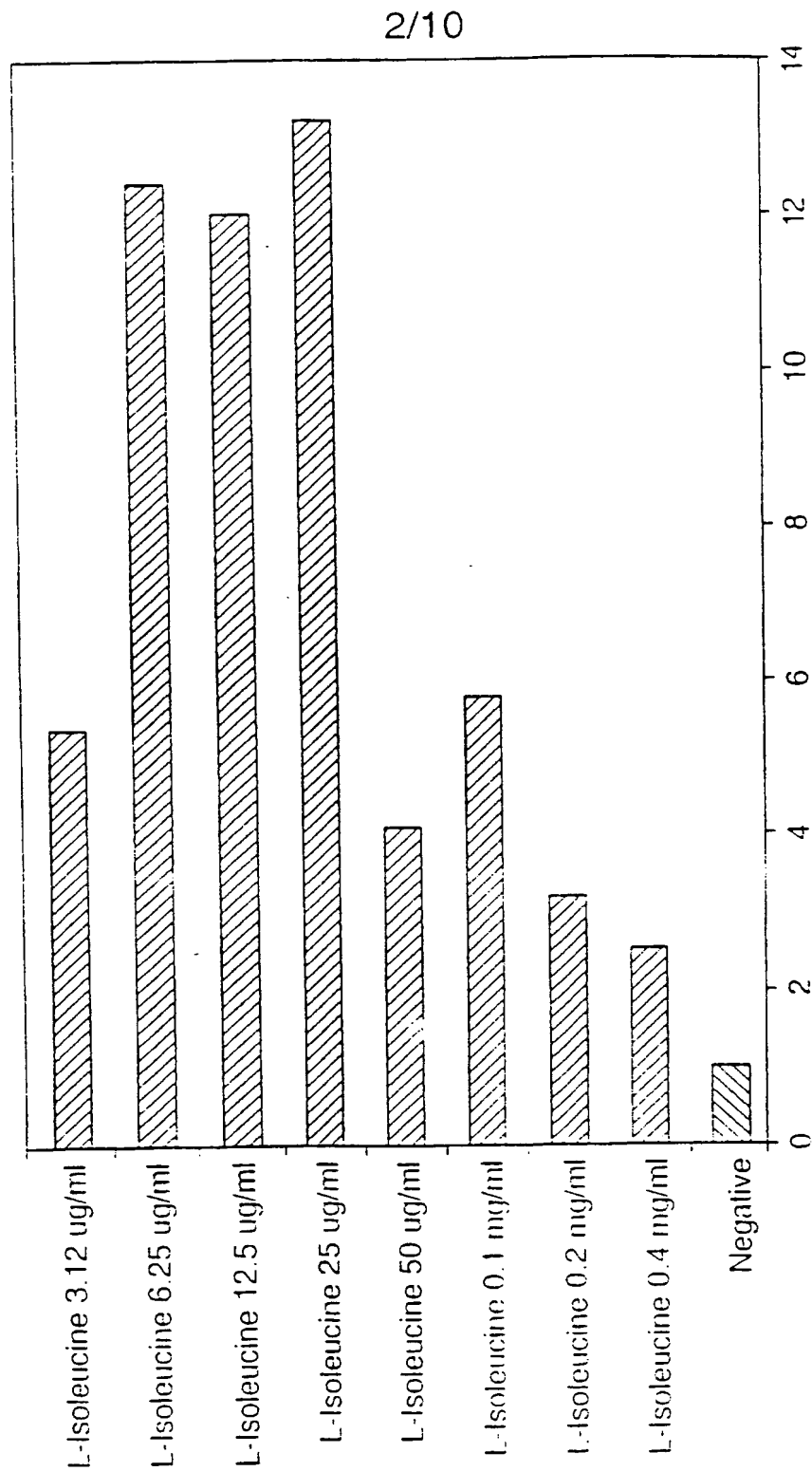
31. A method for improving total weight gain, rate of weight gain, or feed conversion efficiency in livestock comprising administration of an antimicrobial peptide inducing compound.

30

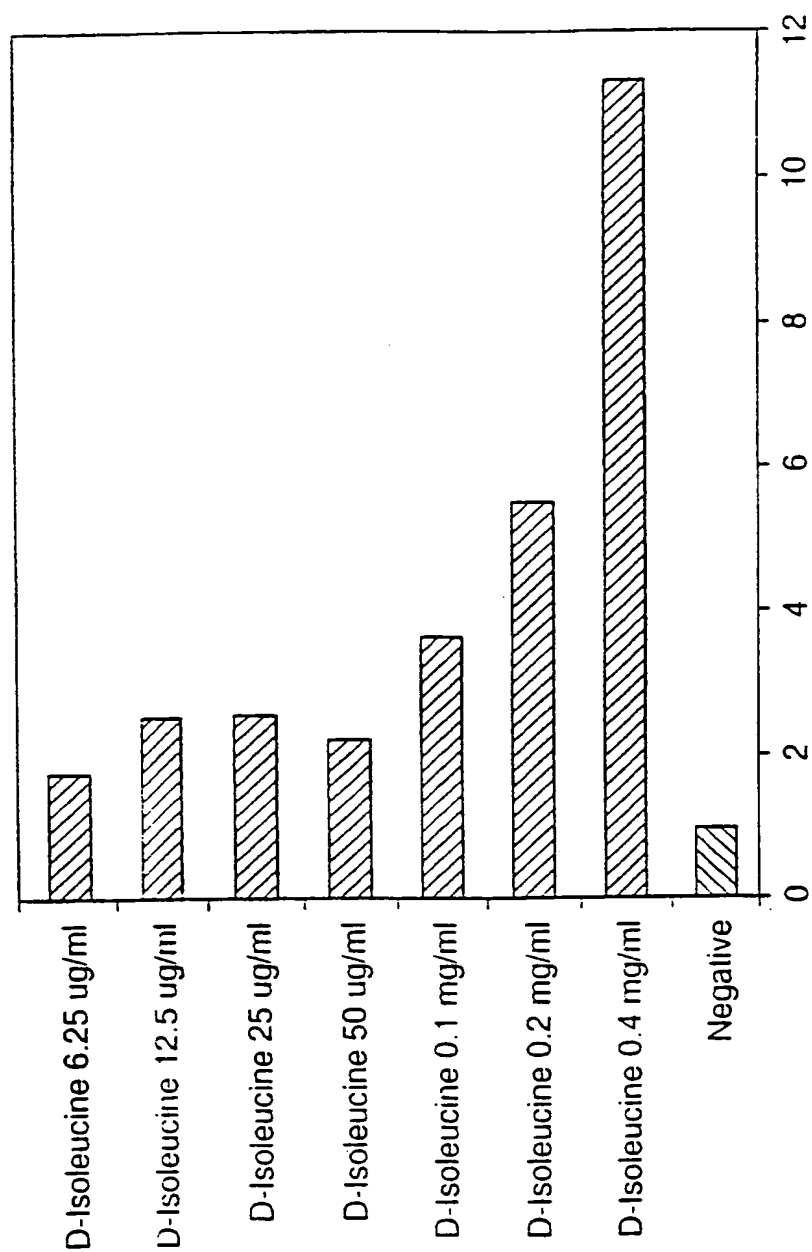
32. A method for improving total weight gain, rate of weight gain, or feed conversion efficiency in livestock comprising administration of isoleucine or active isomers or analogs thereof.

1/10

**FIG. 1**

**FIG. 2**

3/10

**FIG. 3**

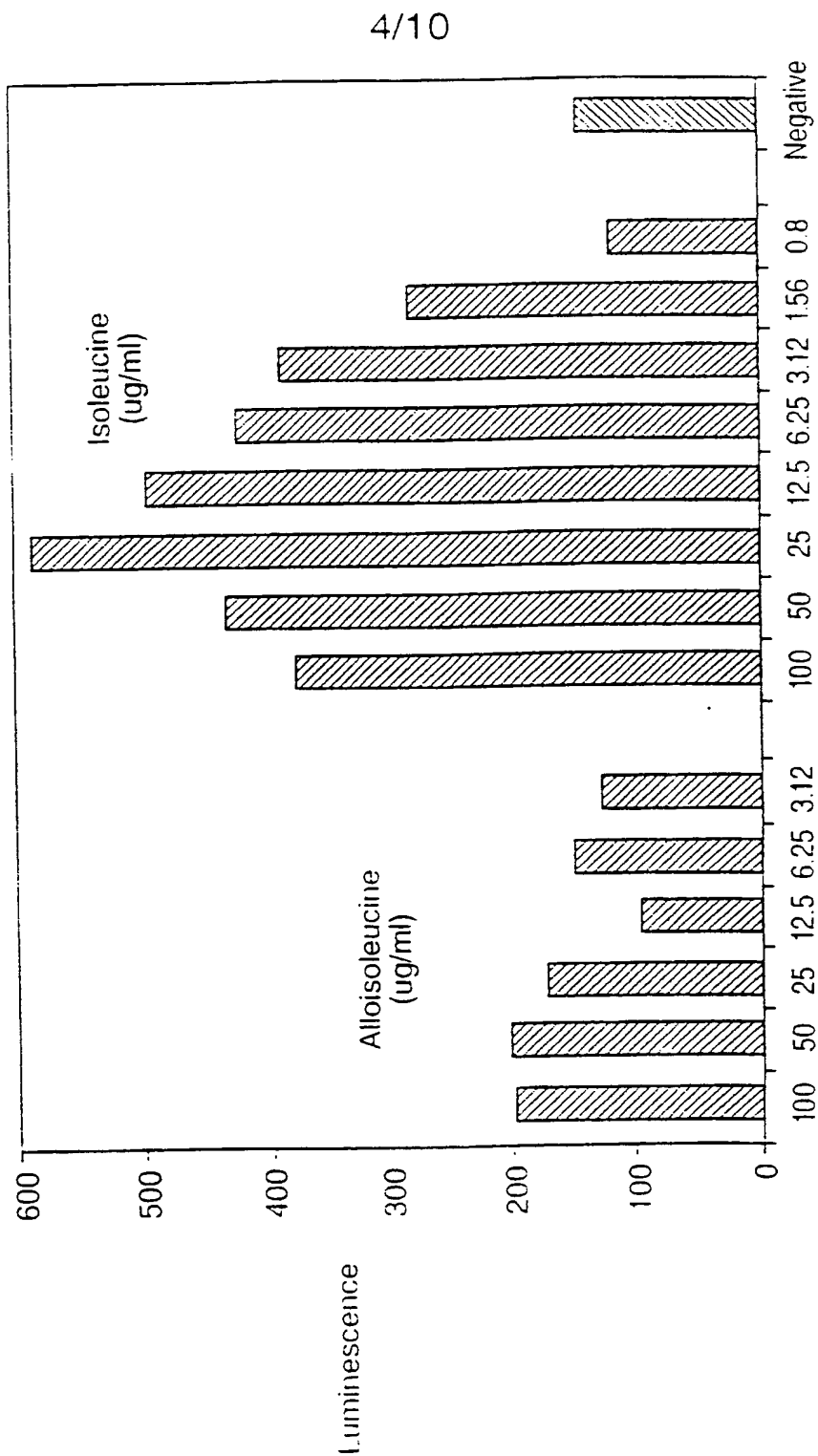
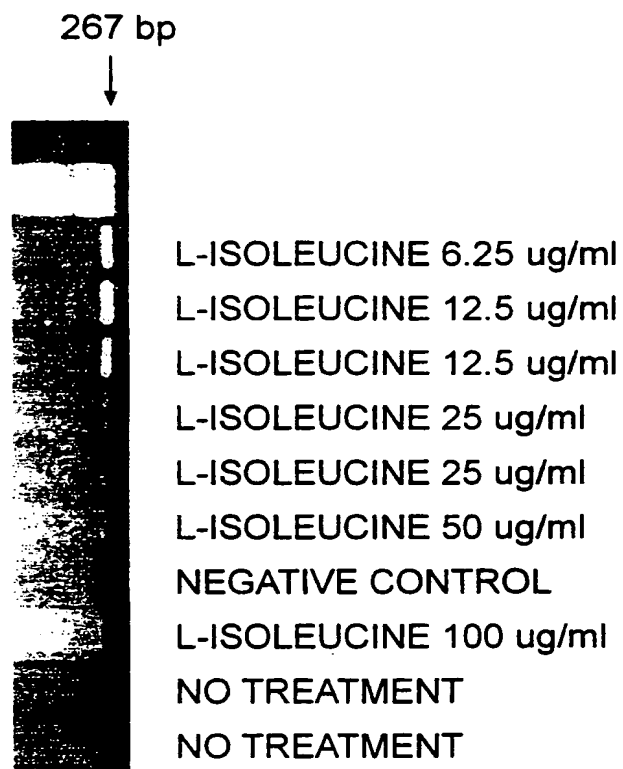


FIG. 4

5/10

**FIG. 5**

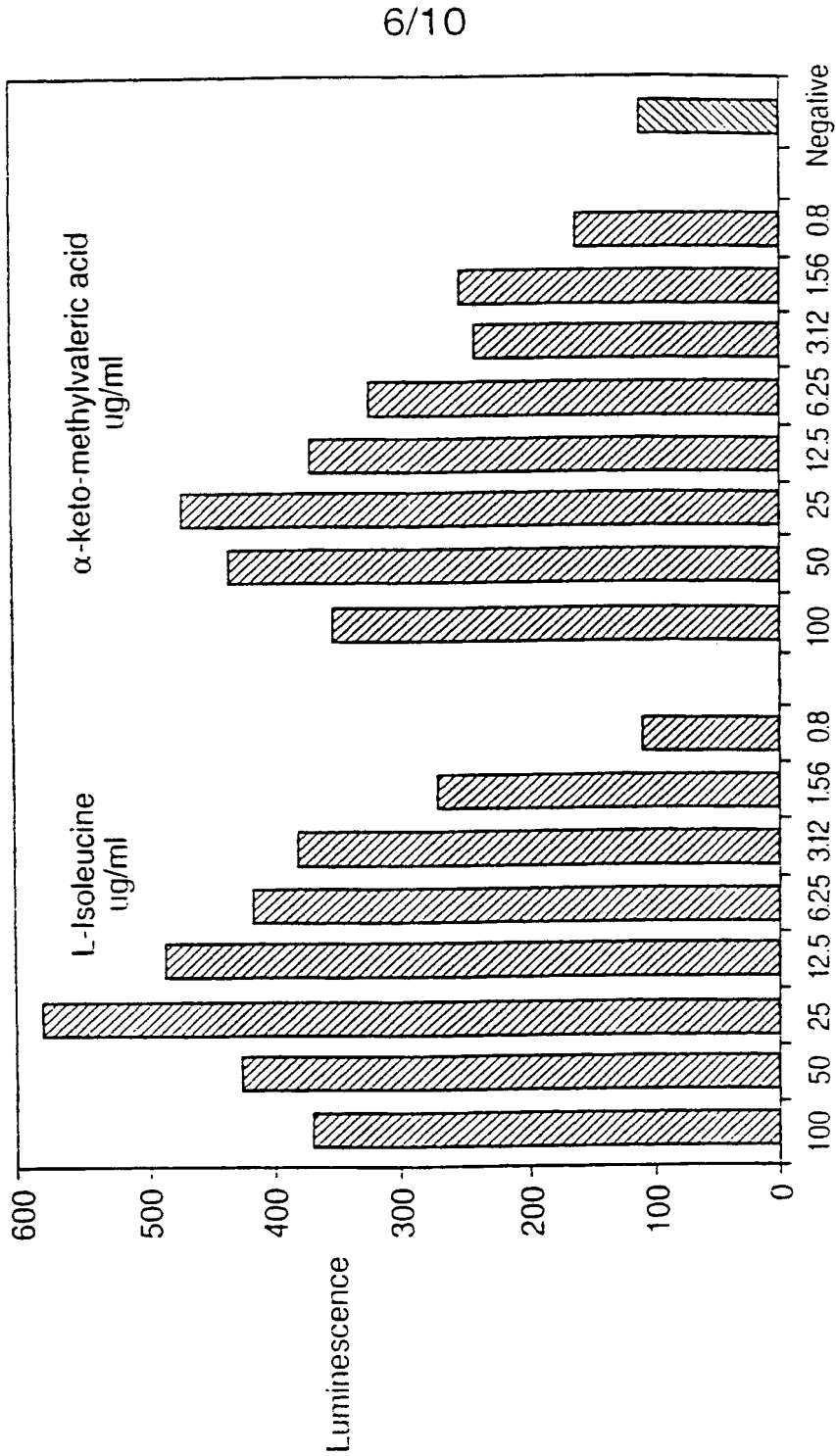


FIG. 6

7/10

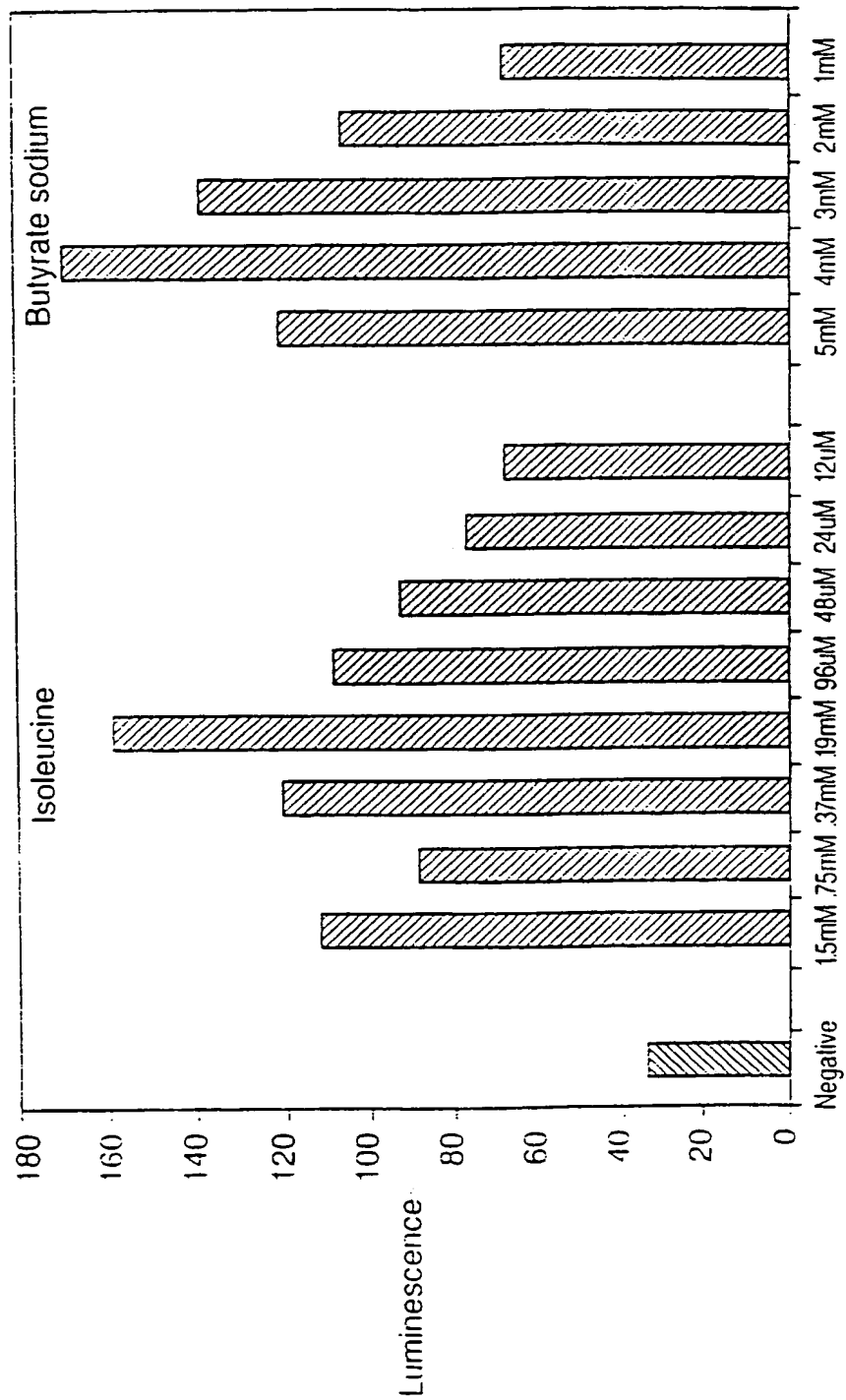


FIG. 7

8/10

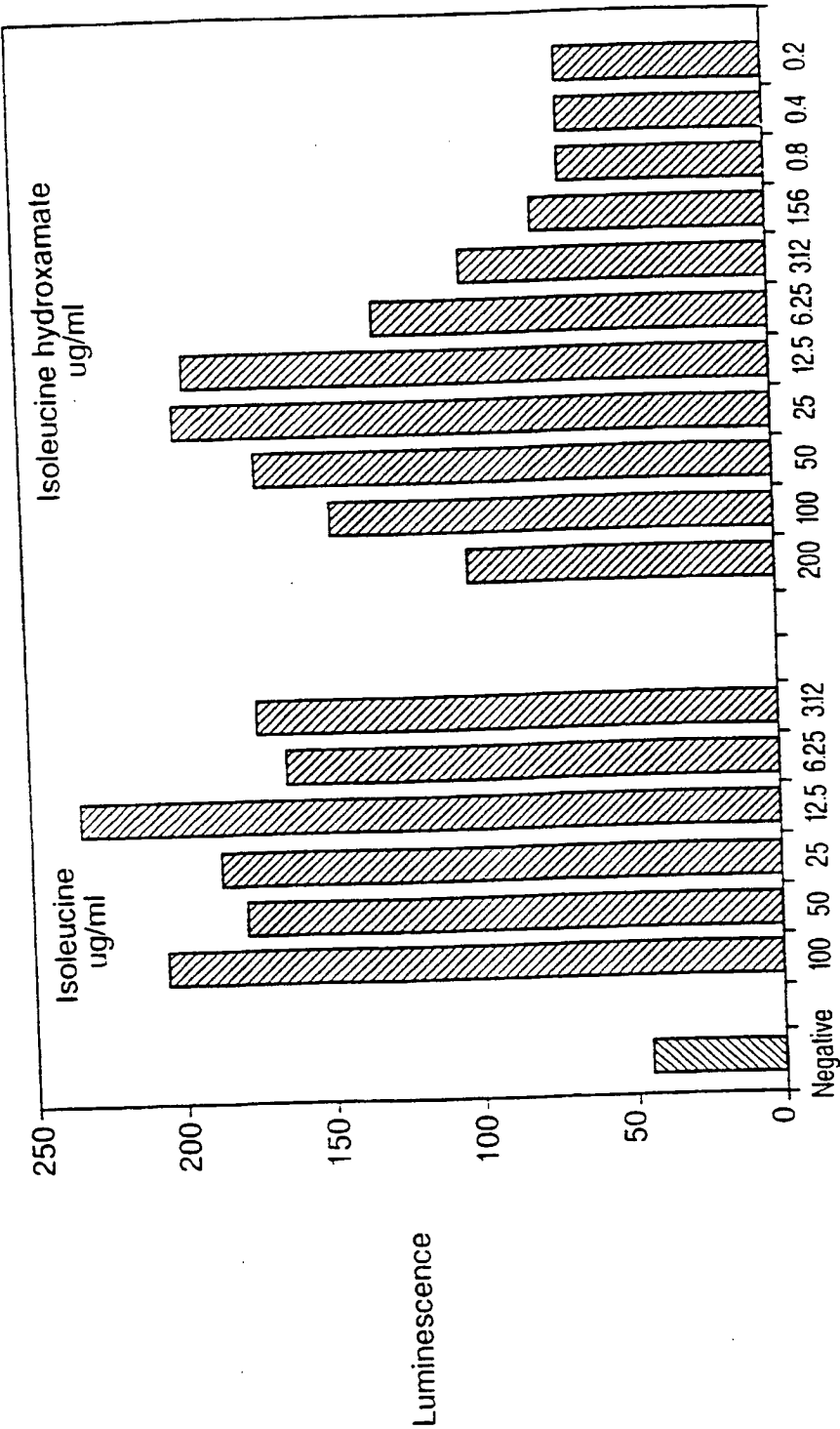
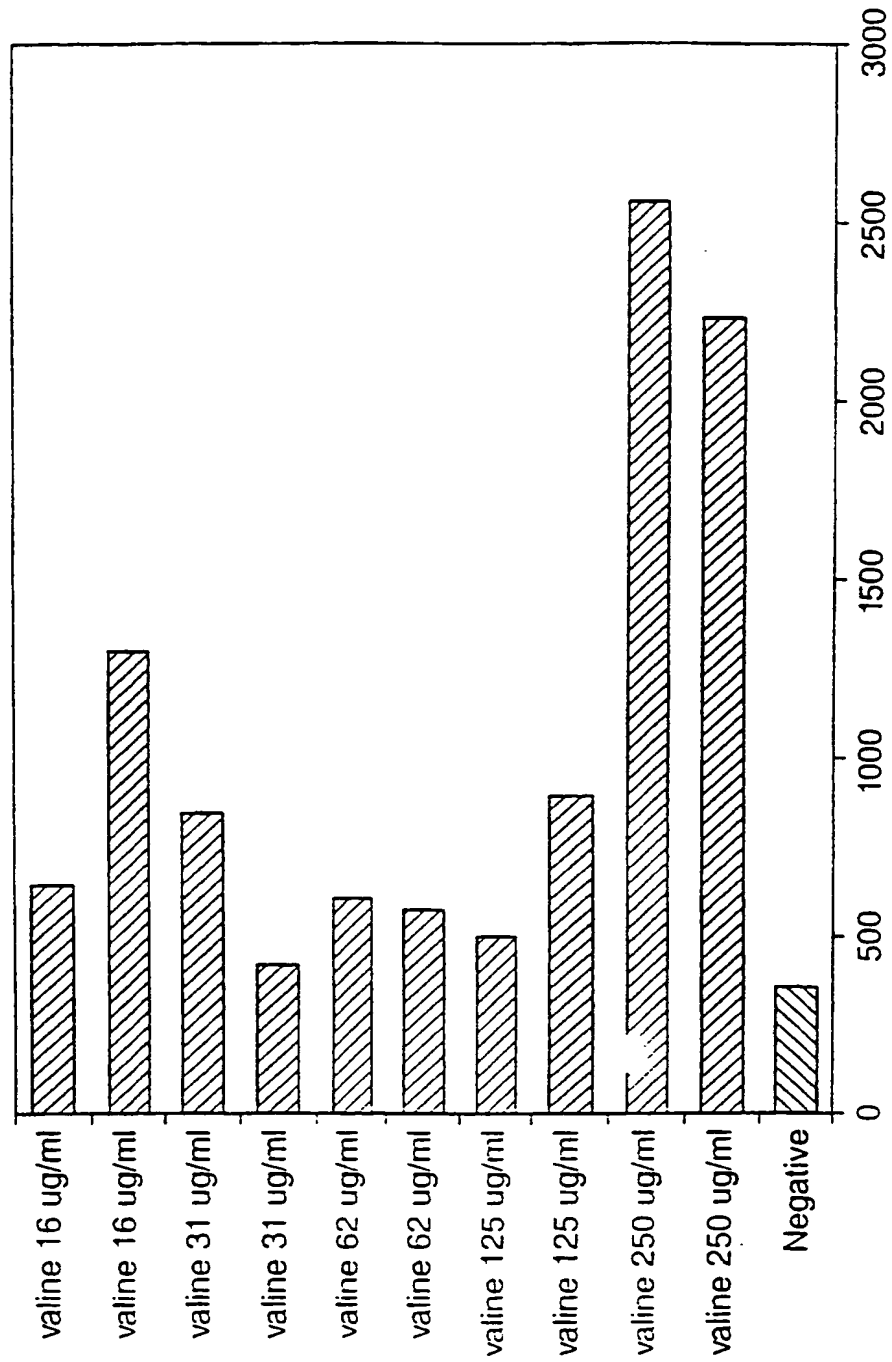


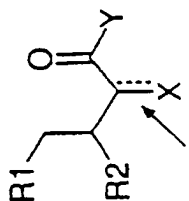
FIG. 8

9/10

**FIG. 9**

10/10

$X = -NH_2, O, H.$
 $Y = -OH, -NHOH, -NH_2, -OCH_3, -OR_3.$
 $R_1 = H, -CH_3, \text{alkyl of 2-5 carbons.}$
 $R_2 = H, -CH_3, \text{alkyl of 2-5 carbons.}$
 $R_3 = \text{alkyl of 2-5 carbons.}$



single or double bond

FIG. 10

INTERNATIONAL SEARCH REPORT

International application No.

PCT/US01/08197

A. CLASSIFICATION OF SUBJECT MATTER

IPC(7) : A61K 31/195, 31/185, 31/19, 31/22.

US CL : 514/12; 530/324, 300; 435/116, 115.

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

U.S. : 514/12; 530/324, 300; 435/116, 115, 6, 252.33, 252.9, 253.5, 325

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)
Please See Continuation Sheet

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO 99/59574 A1 (MAGAININ PHARMACEUTICALS, INC.) 25 November 1999 (25.11.99). See whole document and claims 1-14.	1-14
Y	WO 99/49876 A1 (ABBOTT LABORATORIES) 07 October 1999 (07.10.99). See pages 4-5, 8-10; Table I and II; Figure 2.	1, 4-8
X,P	FEHLBAUM, P. et al. An Essential Amino Acid Induces Epithelial beta-Defensin Expression. Proc. Natl. Acad. Sci. 07 November 2000, Vol. 97, No. 23, pages 12723-12728. See pages 12724-12726; Table 1 and 2; Figure 2.	1-2, 4-9, 11-14
A	SCHRODER, J.-M. Clinical Significance of Epithelial Peptide Antibiotics. BioDrugs May 1999, Vol. 11, No. 5, pages 293-300. See whole document.	1, 4-6
A	O'NEIL, D.A. et al. Expression and Regulation of the Human beta-Defensins hBD-1 and hBD-2 in Intestinal Epithelium. J. Immunol. December 1999, Vol. 163, No. 12, pages 6718-6724. See whole document	1, 4-6

☐ Further documents are listed in the continuation of Box C.

☐ See patent family annex.

* Special categories of cited documents:	
"A" document defining the general state of the art which is not considered to be of particular relevance	"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
"E" earlier application or patent published on or after the international filing date	"X" document of particular relevance: the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
"O" document referring to an oral disclosure, use, exhibition or other means	"&" document member of the same patent family
"P" documents published prior to the international filing date but later than the priority date claimed	

Date of the actual completion of the international search	Date of mailing of the international search report 25 JUN 2001
Name and mailing address of the ISA/US Commissioner of Patents and Trademarks Box PCT Washington, D.C. 20231 Facsimile No. (703)305-3230	Authorized officer Chih-Min Kam Telephone No. (703) 308-0196

Form PCT/ISA/210 (second sheet) (July 1998)

INTERNATIONAL SEARCH REPORT

International application No.

PCT/US01/08197

Box I Observations where certain claims were found unsearchable (Continuation of Item 1 of first sheet)

This international report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☐ Claim Nos.:
because they relate to subject matter not required to be searched by this Authority, namely:
2. ☐ Claim Nos.:
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:
3. ☒ Claim Nos.: 15
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of Item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:
Please See Continuation Sheet

1. ☐ As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☒ No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.: 1-14

Remark on Protest

☐
☐

- The additional search fees were accompanied by the applicant's protest.
- No protest accompanied the payment of additional search fees.

INTERNATIONAL SEARCH REPORT

International application No.

PCT/US01/08197

BOX II. OBSERVATIONS WHERE UNITY OF INVENTION IS LACKING:

This International Search Authority has found 4 inventions claimed in the International Application covered by the claims indicated below:

- I. Claims 1-14, drawn to a method of eliciting the production of defensins in eukaryotic cells comprising exposing the cells to a composition comprising isoleucine or active isomers or analogs.
- II. Claims 16-20, drawn to a method of treating or preventing an infection or other disease state in a patient by administering to patient a composition consisting of isoleucine or active isomers or analogs.
- III. Claims 21-30, drawn to a method of increasing numbers of probiotic or commensal bacteria at an epithelial surface or in an epithelially lined compartment comprising administration of a composition comprising isoleucine or active isomers or analogs, and a composition comprising isoleucine or active isomers or analogs to elicit increases in probiotic or commensal bacteria.
- IV. Claims 31-32, drawn to a method for improving total weight gain, rate of weight gain, or feed conversion efficiency in livestock comprising administration of an antimicrobial peptide inducing compound, or isoleucine or active isomers or analogs.

This international Searching Authority considers that the international application does not comply with the requirements of unity of invention (Rules 13.1, 13.2 and 13.3) for the reasons indicated below:

The inventions listed as Groups I-IV do not relate to a single general inventive concept under PCT Rule 13.1 because, under PCT Rule 13.2, they lack the same or corresponding special technical features for the following reasons: In this instance, the method of eliciting the production of defensins in eukaryotic cells is known in the art, see WO99/59574. Thus, the special technical feature is known and the claimed subject matter does not define a contribution which each of the claimed inventions, considered as a whole, makes over the prior art. the special technical feature of the Group I invention is the specific method of eliciting the production of defensins in eukaryotic cells claimed therein while the special technical feature of the Group II invention is the particular method of treating or preventing an infection or other disease state in a patient claimed therein, the special technical feature of the Group III invention is the particular method of increasing numbers of probiotic or commensal bacteria at an epithelial surface or in an epithelially lined department, and a composition comprising isoleucine or active isomers or analogs claimed therein, the special technical feature of the Group IV invention is the particular method for improving total weight gain, rate of weight gain, or feed conversion efficiency in livestock claimed therein. Since the special technical feature of the Group I invention is not present in the Groups II-IV claims, the special technical features of the Groups II-IV inventions are not present in the Group I claims, the special technical feature of the Groups II inventions are not present in the Group III-IV claims, and the special technical feature of the Group III inventions are not present in the Group IV claims, unity of invention is lacking.

Continuation of B. FIELDS SEARCHED Item3: STN search on caplus, medline, embase, biosis and scisearch; EAST search on USPAT, DDERWENT, EPO, and JPO. Search terms used: defensin, production, producing, eliciting, isoleucine, valine, butyrate, alpha-ketomethyl valerate, isoleucine hydroxamate, eukaryotic cell, mammalian cell, epithelial cell.

This Page Blank (uspto)